1 Claims

2

3 1. A compound of the following Formula 1:

$$\begin{array}{c|c}
R_{10} & R_{11} \\
R_{10} & B \\
R_{13} & C \\
R_{13} & R_{14}
\end{array}$$

4

5 6

wherein

 R_A is a C_2 to C_{30} saturated or unsaturated

8 hydrocarbon chain;

9

10

11

12

13

14

R₁₀, R₁₁, R₁₃, R₁₄ and R₃ each independently represent H, OH, a C₁₋₆ ether, or a saturated or unsaturated hydrocarbon chain which may be substituted with one or more of nitro, halogen, amino, hydroxyl, ketone or aldehyde group;

15

optionally there is a double bond between C₂ and C₃ of the C ring;

18 19

n represents 0 or 1; and

20

 R_B is a C_2 to C_{15} saturated or unsaturated hydrocarbon chain, and where R_B is present, R_A and R_B are both C_2 to C_{12} aliphatic alkyl chains.

1 2. The compound as claimed in Claim 1 wherein at least one of R₁₀, R₁₁ and R₁₃ represents OH.

3

4 3. The compound as claimed in Claim 2 wherein R_{10} and/or R_{11} represents OH.

6

7 4. The compound as claimed in any one of Claims 1 to 3 wherein R_3 , R_{11} and R_{13} all represent OH.

9

10 5. The compound as claimed in any one of Claims 1 11 to 3 wherein R₃, R₁₀ and R₁₃ all represent OH.

12

13 6. The compound as claimed in any one of Claims 1 14 to 5 wherein there is a double bond between C₂ 15 and C₃ of the C ring.

16

7. The compound as claimed in any one of Claims 1 to 6 where the backbone of R_A has eight, nine or ten carbon atoms.

20

21 8. The compound as claimed in any one of Claims 1 to 7 where R_A is attached to position 7 of the 23 A ring of the flavonoid group.

24

25 9. The compound as claimed in any one of Claims 1 to 8 wherein R_A has the following structure:

$$H_3C$$
 CH_3
 CH_2
 m

- 28 wherein
- n is an integer from 1 to 7; and

m is an integer from 1 to 7.

2

3 10. The compound as claimed in any one of Claims 1 4 to 8 wherein R_A has the following structure:

5

The compound as claimed in any one of Claims 1 to 8 wherein R_A has the following structure:

8

9 wherein n is an integer from 2 to 27.

10

11 12. The compound as claimed in any one of Claims 1 12 to 8 wherein R_A has the following structure:

13

14 wherein

x is an integer from 1 to 25;

16

y is an integer from 1 to 25;

18

and wherein x + y = 25 or less.

20

21 13. The compound as claimed in any one of Claims 1 22 to 12 wherein R_A has the following structure:

$$H_3C$$
 CH_3
 CH_2
 CH_2

OH

1 wherein

n is an integer from 1 to 7; and

3

m is an integer from 1 to 7.

5

6 14. The compound as claimed in any one of Claims 1
7 to 13 wherein the flavonoid group has one of
8 the following structures:

9 10

11 12

13 15. The compound as claimed in any one of Claims 1 14 to 14 having one of the following structures:

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1 16. A composition comprising a compound as claimed 2 in any one of Claims 1 to 15 and at least one 3 pharmaceutical excipient or carrier.

4

5 17. The composition as claimed in Claim 16 which is a sunscreen.

7

8 18. A method of preventing UV damage to the skin of
9 a mammalian animal, said method comprising
10 administering a therapeutically effective
11 amount of the composition of Claim 17 to said
12 skin prior to UV exposure.

13

14 19. The method as claimed in Claim 18 wherein said mammalian animal is a human.

1	20.	The method as claimed in either one of Claims
2		18 and 19 wherein said composition is applied
3		topically to said skin.
4		
5	21.	The composition as claimed in Claim 16 which is
6		a skincare composition.
7		
8	22.	The composition as claimed in Claim 21 further
9		containing emollients and moisturisers.
10		
L1	23.	The composition as claimed in either one of
12		Claims 21 and 22 for preventing or reversing
13		the effects of ageing, of reducing apparent
14		wrinkling and/or treating or preventing dry
1.5		skin.
16		
17	24.	A foodstuff stabiliser composition comprising a
18		compound as claimed in any one of Claims 1 to
19		15.
20		
21	25.	The composition as claimed in Claim 24 in the
22		form of an emulsion having a low fat:high water
23		content.
24		
25	26.	A method of treating a patient having a disease
26		or disorder involving oxidative damage, said
27		method comprising the step of administering a
28		therapeutically effective amount of the
29		composition of Claim 16 to said patient.
30		

1	27.	The method as claimed in Claim 26 wherein said
2		patient is a human.
3		
4	28.	The method as claimed in either one of Claims
5		26 and 27 wherein the disease or disorder
6		involving oxidative damage is selected from the
7		group consisting of cancer, heart disease,
8		neurological disorders, auto-immune disorders,
9		ischaemia-reperfusion injury, diabetic
LO		complications, septic shock, hepatitis,
L1		atherosclerosis and complications arising from
L 2		HIV or Hepatitis B.
L3		·
L4	29.	The method as claimed in Claim 28 wherein the
L5		disease or disorder is an ischaemia-reperfusion
L6		injury or Alzheimer's disease.
L7		
L8	30.	A prophylactic method of treatment to prevent
L9		or reduce the severity of a disease or disorder
20		involving oxidative damage in the tissues of a
21		patient, said method comprising the step of
22		administering a therapeutically effective
23		amount of the composition of Claim 16 to said
24		patient.
25		
26	31.	The method as claimed in Claim 30 wherein said
27		patient is a human.
28		
29	32.	The method as claimed is either one of Claims
30		30 and 31 wherein the disease or disorder
31		involving oxidative damage is selected from the

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1		group consisting of cancer, heart disease,
2		neurological disorders, auto-immune disorders,
3		ischaemia-reperfusion injury, diabetic
4		complications, septic shock, hepatitis,
5		atherosclerosis and complications arising from
6		HIV or Hepatitis B.
7		
8	33.	The method as claimed in Claim 32 wherein the
9		disease or disorder is an ischaemia-reperfusion
10		injury or Alzheimer's disease.
11		
12	34.	The use of a compound of Formula 1 as claimed
13		in any one of Claims 1 to 15 for the
14		manufacture of a medicament for the treatment
15		of a disease or disorder involving oxidative
		_
16		damage.
16 17		damage.
	35.	The use as claimed in Claim 34 wherein the
17	35.	
17 18	35.	The use as claimed in Claim 34 wherein the
17 18 19	35.	The use as claimed in Claim 34 wherein the disease or disorder is cancer, heart disease,
17 18 19 20	35.	The use as claimed in Claim 34 wherein the disease or disorder is cancer, heart disease, neurological disorders, auto-immune disorders,
17 18 19 20 21	35.	The use as claimed in Claim 34 wherein the disease or disorder is cancer, heart disease, neurological disorders, auto-immune disorders, ischaemia-reperfusion injury, diabetic
17 18 19 20 21	35.	The use as claimed in Claim 34 wherein the disease or disorder is cancer, heart disease, neurological disorders, auto-immune disorders, ischaemia-reperfusion injury, diabetic complications, septic shock, hepatitis,
17 18 19 20 21 22	35.	The use as claimed in Claim 34 wherein the disease or disorder is cancer, heart disease, neurological disorders, auto-immune disorders, ischaemia-reperfusion injury, diabetic complications, septic shock, hepatitis, atherosclerosis, and complications arising from
17 18 19 20 21 22 23		The use as claimed in Claim 34 wherein the disease or disorder is cancer, heart disease, neurological disorders, auto-immune disorders, ischaemia-reperfusion injury, diabetic complications, septic shock, hepatitis, atherosclerosis, and complications arising from
17 18 19 20 21 22 23 24		The use as claimed in Claim 34 wherein the disease or disorder is cancer, heart disease, neurological disorders, auto-immune disorders, ischaemia-reperfusion injury, diabetic complications, septic shock, hepatitis, atherosclerosis, and complications arising from an immune response to HIV or Hepatitis B.
17 18 19 20 21 22 23 24 25		The use as claimed in Claim 34 wherein the disease or disorder is cancer, heart disease, neurological disorders, auto-immune disorders, ischaemia-reperfusion injury, diabetic complications, septic shock, hepatitis, atherosclerosis, and complications arising from an immune response to HIV or Hepatitis B. A method of manufacturing a compound of Formula
17 18 19 20 21 22 23 24 25 26		The use as claimed in Claim 34 wherein the disease or disorder is cancer, heart disease, neurological disorders, auto-immune disorders, ischaemia-reperfusion injury, diabetic complications, septic shock, hepatitis, atherosclerosis, and complications arising from an immune response to HIV or Hepatitis B. A method of manufacturing a compound of Formula 1 as claimed in any one of Claims 1 to 15, said
17 18 19 20 21 22 23 24 25 26 27		The use as claimed in Claim 34 wherein the disease or disorder is cancer, heart disease, neurological disorders, auto-immune disorders, ischaemia-reperfusion injury, diabetic complications, septic shock, hepatitis, atherosclerosis, and complications arising from an immune response to HIV or Hepatitis B. A method of manufacturing a compound of Formula 1 as claimed in any one of Claims 1 to 15, said method comprising providing an intermediate

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group where the metal is directly attached to R_A, and R_A is a C₂ to C₃₀ saturated or unsaturated alkyl chain, and R_AM is capable of participating in transition metal catalysed cross-coupling reactions; and intermediate compound B has the following structure:

$$(X)_{m}$$
 $(X)_{m}$
 $(X)_$

wherein

R₁₂ represents OH or an O-protecting group
R₃, R₁₀, R₁₁, R₁₃, and R₁₄ each independently
represent H, OH, C₁ to C₄ aliphatic alkyl group
or an O-protecting group where required, and
optionally there is a double bond between C₂
and C₃ of the C ring;
X is a halogen, O-trifluoromethane sulphonate
or any other group used in cross-coupling
reactions; and
m = 1 or 2,

and reacting intermediate compound A with intermediate compound B by transition metal catalysed cross-coupling reactions and subsequently deprotecting at least one OH group.

1 37. A method as claimed in Claim 36 wherein R_AM is 2 an organomagnesium, organozinc, organoboron or 3 organotin compound.

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The method as claimed in either one of Claims
and 37 wherein the catalyst is a palladium,
nickel or iron complex.

8

A method of manufacturing a compound of Formula 9 39. 1 as claimed in any one of Claims 1 to 15, said 10 method comprising providing an intermediate 11 Compound C and an intermediate Compound D, 12 wherein said intermediate Compound C has the 13 structure RACHCHR wherein RA is as defined in 14 Formula 1, and wherein intermediate Compound D 15 has a structure: 16

17 18

19 40. The method as claimed in Claim 39 wherein the catalyst is:

21

22

1 41. A method of manufacturing a compound of Formula 2 1 as claimed in any one of Claims 1 to 15, said 3 method comprising providing an intermediate 4 Compound E of formula:

$$R_{A_{n_{N_{n}}}}$$
 or $R_{A_{n_{N_{n}}}}$ $O-R_{C}$

5

and constructing a flavonol core on said intermediate Compound E.

9

10 42. The method as claimed in Claim 41 wherein said 11 flavonol core is formed by Algar-Flynn-Oyamada 12 (AFO) oxidation.

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14 43. The method as claimed in Claim 41 wherein said 15 flavanol core is formed by Baker-Verkataraman 16 rearrangement.

17

18 44. The method as claimed in any one of Claims 40
19 to 43 wherein said intermediate Compound E is
20 formed by a transition metal catalysed cross21 coupling reaction.

22

23 45. The method as claimed in any one of Claims 40 24 to 43 wherein said intermediate Compound E is 25 formed by alkene cross-metathesis.